

Muon spin relaxation and neutron scattering investigations of the noncentrosymmetric heavy-fermion antiferromagnet CeRhGe_3

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The magnetic ground state of CeRhGe_3 has been investigated using magnetic susceptibility, heat capacity, neutron diffraction, muon spin relaxation (μSR), and inelastic neutron scattering (INS) techniques. Our μSR study clearly reveals the presence of two frequencies below $T_{\text{N}2} = 7$ K and three frequencies between 7 K and $T_{\text{N}1} = 14.5$ K, indicating long-range magnetic ordering of the Ce^{3+} moment. The temperature dependence of the highest frequency follows a mean-field order parameter. Our powder neutron diffraction study at 1.5 K reveals the presence of magnetic Bragg peaks, indexed by the propagation vector $\mathbf{k} = (0, 0, 3/4)$ with the Ce^{3+} magnetic moment $\sim 0.45(9) \mu_{\text{B}}$ along the c axis. INS studies at 18 K (i.e., above $T_{\text{N}1}$) show the presence of two well-defined crystal-field (CEF) excitations at 7.5 and 18 meV. At 10 and 4.5 K, a very small increase has been observed in the CEF excitation energies. At 100 K, both CEF excitations broaden and a broad quasielastic component has also been observed. Further, the low-energy INS study reveals the presence of a nearly temperature-independent quasielastic linewidth between 16 and 60 K, which indicates a Kondo temperature $T_{\text{K}} = 12.6(3)$ K. The presence of well-defined CEF excitations in CeRhGe_3 suggests local moment magnetism and may explain the absence of pressure-induced superconductivity. Analyzing the INS data based on a CEF model, we have evaluated the CEF ground-state wave functions and ground-state moment. The observed small value of the ordered moment along the c axis, deduced from the neutron diffraction data, contrasts with the ab -plane moment direction predicted by the single-ion CEF anisotropy and indicates the presence of two-ion anisotropic magnetic exchange interactions, which govern the direction of the moment.

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I. INTRODUCTION

Strongly correlated electron systems, displaying heavy-fermion (HF) behavior, have attracted considerable attention both in theoretical and experimental condensed matter physics due to the many exotic properties they can exhibit.¹⁻⁴ One of the most attractive and elusive phenomena of these HF systems is the coexistence of magnetism and superconductivity (SC), which is at the forefront of condensed matter research. The SC in HF systems cannot be explained by the Bardeen-Cooper-Schrieffer (BCS) theory relevant for conventional s -wave phonon-mediated superconductors. Many HF compounds, having centrosymmetric crystal structures, exhibit unconventional SC close to a quantum critical point (QCP), where magnetic order disappears and SC emerges. The region where SC exists can be reached as a function of pressure, magnetic field, or alloying, implying that magnetic fluctuations are playing an important role in the formation of the Cooper pairs.³ In centrosymmetric materials, the spin and orbital parts of the pair wave function can be treated individually because the conduction bands are degenerate even if the spin-orbit coupling is strong.² Therefore, one can express the Cooper pair as either a spin-singlet or a spin-triplet pairing. The pairing mechanism in heavy-fermion superconductors (HFSC) is not yet understood, as is also the case in the high T_c cuprates (HTSC) and the newly discovered Fe-based superconductors.^{3,4} One of the current hypotheses for these unconventional superconductors is that the superconducting pairing is mediated by magnetic interactions, the so-called

“magnetic glue.”^{3,5} Despite large volumes of data available in the literature on HTSC and HFSC, the origin of the SC and pairing mechanism in these classes of materials remains difficult to understand.

Recently, we have been investigating the physical properties of RTX_3 (R = rare earth, T = transition metal, X = Si, Ge, Sn) compounds using μSR and neutron scattering techniques.⁶⁻⁹ Ce compounds of this series, CeTX_3 (X = Si and Ge), belong to the family of recently discovered noncentrosymmetric HFSC whose superconducting ground-state properties are dictated by an antisymmetric spin-orbit coupling (ASOC) as a consequence of the lack of inversion symmetry, along the $[0\ 0\ 1]$ direction, in the tetragonal crystal structure.¹⁰⁻¹⁹ This ASOC, in turn, results in the formation of a mixed pairing wave function with spin-singlet and spin-triplet components in the superconducting state. The relationship between SC and the lack of inversion symmetry is therefore a central issue, which needs to be clarified both theoretically and experimentally in f -electron systems. Among these systems, CeTX_3 compounds exhibit many interesting properties (see Table I). For example, at ambient pressure CeCoSi_3 shows SC at 0.8-1.3 K,^{10,11} while CeRhSi_3 and CeIrSi_3 order antiferromagnetically (AFM) at 1.5 and 5 K, respectively,¹²⁻¹⁵ and also reveal SC over a broad range of applied pressures below 1.1 and 1.6 K, respectively.¹²⁻¹⁵

The Ge-based compounds CeTGe_3 also exhibit interesting magnetic and superconducting properties.¹⁶⁻¹⁸ CeRhGe_3 and CeIrGe_3 exhibit three successive magnetic phase transitions,

TABLE I. A summary of the physical properties of CeTX₃ compounds crystallizing in the BaNiSn₃-type noncentrosymmetric tetragonal crystal structure (*I4mm*, No. 107): the magnetic ground state, superconducting transition temperature (T_c) at pressure (P), paramagnetic Curie Weiss temperature (θ_p), electronic specific-heat coefficient (γ), Kondo temperature (T_K), and residual resistivity (ρ_0). The magnetic ground states are antiferromagnetic (AFM) with Néel temperature T_N , paramagnetic (PM), and spin glass (SG). VF and HF denote valence fluctuation and HF states, respectively.

Compound magnetism	θ_p (K)	T_N (K)	γ mJ/mol K ²	T_K (K)	T_c (K)	P (kbar)	ρ_0 ($\mu\Omega$ cm)	Ref.
CeCoSi ₃ PM (VF)	-840		37	900	0.7-1.4	0	12	10,11,54
CeRuSi ₃ PM (VF)							~28	10
CeRhSi ₃ AFM (HF)	-128	1.6	120	100	1	23.1	~6	12,16
CePdSi ₃ AFM	26	3/5.2	57				~2	55
CeOsSi ₃ PM (VF)							~20	10
CeIrSi ₃ AFM (HF)	-142	5.0	125	100	1.6	25	~10	14,16
CePtSi ₃ AFM	-211/12	4.8/2.4	29				16.6	19
CeFeGe ₃ PM (HF)	-90		150	100			~8	56
CeCoGe ₃ AFM	-71/-29	21/12/8 32					1/2.5	20
CeRhGe ₃ AFM	-28	14.6/10/0.55	40				~2(10 ^a)	16
CeIrGe ₃ AFM	-21	8.7/4.7/0.7	80				~5	16
^b CeNiGe ₃ AFM	-17	5.5	45	4.3	0.48	65	~8	22,57
^c CeRuGe ₃ SG (HF)	-7.5		428					58

^aPresent work.

^bCrystallizes in the orthorhombic SmNiGe₃-type structure (*Cmmm*, No. 65).

^cCrystallizes in the cubic structure (*Pm3n*, No. 223).

$T_{N1} = 14.6$ K, $T_{N2} = 7$ K, $T_{N3} = 0.55$ K for CeRhGe₃ and $T_{N1} = 8.7$ K, $T_{N2} = 4.7$ K, $T_N = 0.7$ K for CeIrGe₃, respectively.^{16,17} T_{N1} for CeRhGe₃ increases with increasing pressure and reaches a maximum of 21.3 K at 8.0 GPa.¹⁷ Whereas for CeIrGe₃, T_{N1} remains nearly constant, but T_{N2} increases with pressure and merges with T_{N1} at 4 GPa.¹⁷ Upon increasing the pressure still further, SC has been observed in CeIrGe₃ above 20 GPa with a transition temperature of 1.6 K at 24 GPa.¹⁸ Interestingly, CeCoGe₃ also exhibits unusual magnetic properties with an ordering temperature as high as 21 K²⁰ and pressure-induced SC ($T_{sc} = 0.7$ K at 5.5 GPa).²¹ Furthermore, the Kondo lattice system CeNiGe₃, which crystallizes in the orthorhombic space group (*Cmmm*), orders antiferromagnetically below 5.5 K and exhibits pressure-induced HFSC, showing two domes of SC with a maximum $T_{sc} = 0.3$ K around 3.5 GPa and 0.4 K around 7 GPa and Fermi liquid behavior at higher pressure.^{22,23}

To gain insight into the low-temperature physical properties, anisotropy, and strength of hybridization in CeRhGe₃, the magnetic and thermal properties have been investigated using various experimental techniques. The aim of the current study is to provide a direct comparison of this compound with CeRhSi₃, CeIrSi₃, and CeIrGe₃, which exhibit noncentrosymmetric HFSC under the application of pressure. To develop any realistic theoretical model of magnetism and SC, information on the magnetic structure and CEF ground-state wave functions is important. Our μ SR and neutron diffraction studies clearly reveal long-range magnetic ordering in CeRhGe₃ below 14.5 K. Inelastic neutron scattering (INS) reveals two well-defined crystal-field (CEF) excitations at 7.5 and 18 meV, indicating the localized nature of the 4f-electrons in CeRhGe₃.

II. EXPERIMENTAL DETAILS

Polycrystalline samples (mass ~7 g) of CeRhGe₃ and its nonmagnetic phonon reference LaRhGe₃ were prepared by the standard arc-melting method starting with a stoichiometric mixture of the high-purity elements (Ce, La: 99.9%, Rh: 99.99%, Ge: 99.999%) on a water-cooled copper hearth under an inert argon atmosphere. To improve the homogeneity and phase purity, the samples were flipped and remelted several times during the preparation process. The as-cast samples were wrapped in tantalum foils and annealed for 1 week at 950 °C under a dynamic vacuum to improve the phase formation. Magnetic susceptibility was measured using a commercial superconducting quantum interference device (SQUID) magnetometer (MPMS, Quantum Design). Specific heat measurements were performed by the relaxation method in a physical properties measurement system (PPMS, Quantum Design).

The neutron scattering and muon spin relaxation (μ SR) experiments were carried out at the ISIS Pulsed Neutron and Muon Facility of the Rutherford Appleton Laboratory, United Kingdom. The neutron diffraction measurements were carried out on the general materials (GEM) (at 300 K to check the phase purity) and WISH²⁴ (at 1.5 and 20 K to determine the magnetic structure) time-of-flight diffractometers. The powdered sample was put into a vanadium can with a diameter of 6 mm and placed inside a standard helium cryostat on WISH, and the measuring time was 6 h at each temperature. The INS measurements were carried out on the time-of-flight (TOF) MARI spectrometer between 4.5 and 100 K. The powder samples were wrapped in a thin Al foil and mounted inside a thin-walled cylindrical Al can, which was cooled down to 4.5 K inside a top-loading closed-cycle refrigerator with He-exchange gas around the samples. The measurements were

performed with an incident neutron energy $E_i = 45$ meV, corresponding to a full width at half maximum (FWHM) instrumental resolution of 1.66 and 1.0 meV at zero and 20 meV energy transfer, respectively. Additional low-energy neutron scattering measurements were carried out using the TOF spectrometer IN6 at the Institut Laue-Langevin (ILL), Grenoble, France with $E_i = 3.1$ meV (elastic energy resolution 0.07 meV) and 4.86 meV (elastic energy resolution 0.17 meV). The μ SR experiments were performed on the MuSR spectrometer in the longitudinal geometry configuration. At ISIS, pulses of muons are implanted into the sample at 50 Hz and with a FWHM of ~ 70 ns. These implanted muons decay with a half-life of $2.2 \mu\text{s}$ into positrons, which are emitted preferentially in the direction of the muon spin axis. Each positron is time stamped, and therefore the muon polarization can be followed as a function of time. The MuSR spectrometer comprises 64 detectors. The detectors before the sample (F) are summed together as well as the detectors after the sample (B). The muon polarization then can be determined by

$$G_z(t) = \frac{N_F(t) - \alpha N_B(t)}{N_F(t) + \alpha N_B(t)}, \quad (1)$$

where N_F and N_B are the counts in the forward and backward detectors, respectively, and α is a calibration constant. The sample was mounted on an Ag plate and covered with a thin layer of Mylar. The sample mount was then inserted into an Oxford Instruments Variox cryostat with a temperature range of 1.2 to 300 K. Any Ag exposed to the muon beam would give a time-independent background.

III. RESULTS AND DISCUSSION

A. Neutron diffraction: Characterization at room temperature

Neutron powder diffraction patterns of both samples at 300 K were collected on the GEM diffractometer, and the analysis was carried out using the GSAS program on the basis of a tetragonal BaNiSn_3 -type structure (space group $I4mm$, No. 107). In this structure, the Ce (or La) and Rh atoms occupy the 2a sites (0 0 z), where $z = z_{\text{Ce}}$ (or z_{La}), z_{Rh} , Ge1

atoms occupy the 2a (0 0 0), while the Ge2 atoms occupy the 4b ($0 \frac{1}{2} z_{\text{Ge2}}$) sites.

A summary of our results is given in Table II, and these values are in agreement with the published data.^{17,25} Our analysis showed that both the samples were nearly single phase. Additionally, when the site occupancy of the Ce and Ge atoms was varied, while keeping the site occupancy of the Rh atom fixed to 100%, we found no improvement in the weighted profile reliability factor (R_{WP}), but the site occupancies of Ce (La) and Ge were found to be about 92 and 98%. As the R_{WP} did not improve, we kept all occupancies fixed to 100% in the final refinement. The nearest-neighbor distances are (in \AA) 4.3960 for Ce-Ce, 3.4226 for Ce-Rh, 3.2104 for Ce-Ge1, 3.2498 for Ce-Ge2, and 2.3900 for Rh-Ge1. The distance between Rh-Ge1 is the shortest among these distances. Further, the four Ge1 atoms are at the shortest distance from the Ce, which suggests the importance of Ce-4f and Ge-4p hybridization on the physical properties of CeRhGe_3 and could explain why the superconducting properties of the Si-based compounds, CeTSi_3 (T = Co, Rh and Ir), are different from those of the Ge-based compounds.

B. Magnetic susceptibility

The dc susceptibility of CeRhGe_3 was measured between 2 and 300 K in an applied magnetic field of 0.1 T with the results shown in Fig. 1(a). The low-temperature susceptibility [see the inset in Fig. 1(a)] shows an anomaly near 14.5 K and a peak at 8.5 K, indicating two magnetic phase transitions in agreement with previous reports.¹⁷ The slight upturn below 3 K is due to another (third) magnetic transition at 0.55 K, which is also seen by Muro *et al.* (see the inset of Fig. 5 in Ref. 16). For temperatures above 150 K, the inverse susceptibility follows a Curie-Weiss (CW) behavior, with an effective paramagnetic moment, $\mu_{\text{eff}} = 2.30(1) \mu_{\text{B}}$ and a CW temperature $\theta_{\text{p}} = +3.1(1)$ K. The small, but positive, value of θ_{p} may be due to either weak ferromagnetic correlations between the Ce ions in the high-temperature range or the presence of CEF and anisotropic two-ion exchange. Support for the latter hypothesis comes from our analysis of the single-crystal susceptibility

TABLE II. A summary of the results obtained from the refinement of the room-temperature neutron powder diffraction data: the lattice parameters, atomic positions, occupancy, and thermal parameters. The site occupancy for all the atoms was fixed to 100%. The reliability factors (weighted profile factor R_{wp}) were calculated by comparing the fit to the data.

	CeRhGe_3	LaRhGe_3		
Space group	$I4mm$	$I4mm$		
a (\AA)	4.39604(2)	4.41978(2)		
c (\AA)	10.02243(7)	10.04588(7)		
R_{wp} (%)	5.05	4.71		
	x	y	z	$U (\times 10^{-2}) \text{\AA}^2$
Ce	0	0	0.58006(7)	0.887(16)
Rh	0	0	0.23847(5)	0.049(13)
Ge1	0	0	0	0.701(11)
Ge2	0	0.5	0.34121(4)	0.817(9)
La	0	0	0.58138(7)	0.681(12)
Rh	0	0	0.23949(9)	0.012(10)
Ge1	0	0	0	0.626(11)
Ge2	0	0.5	0.34204(6)	0.720(9)

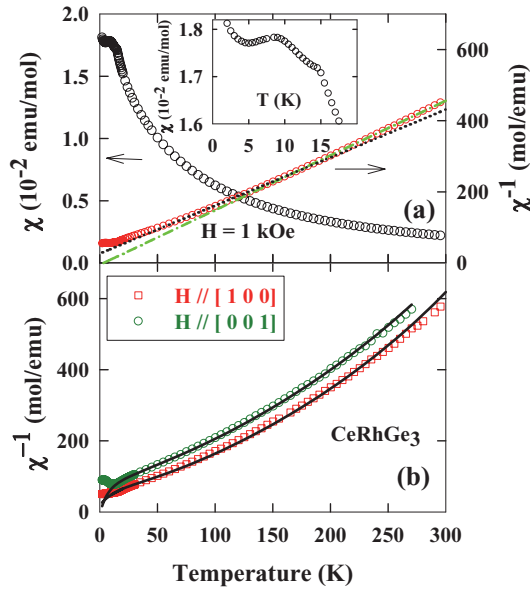


FIG. 1. (Color online) (a) Temperature dependence of the magnetic susceptibility (left y axis) and the inverse magnetic susceptibility (right y axis) of polycrystalline CeRhGe₃. The dotted and dash-dotted lines show fits to CW behavior in the low- and high-temperature regions, respectively. The inset figure shows the low-temperature behavior of the susceptibility. (b) Temperature dependence of the inverse magnetic susceptibility of CeRhGe₃ single crystal from Ref. 17. The solid lines show the fit based on the crystal-field model (together with INS data) including anisotropic molecular field parameters (λ^ξ) and temperature-independent constant susceptibility (χ_0^ξ). The form of the susceptibility used in the direction ξ ($= \perp$ and \parallel to c axis) is given by $\chi^\xi = \chi_{\text{CEF}}^\xi / (1 - \lambda^\xi \chi_{\text{CEF}}^\xi) + \chi_0^\xi$, where χ_{CEF}^ξ is the single-ion susceptibility calculated using the CEF model. The values of the parameters obtained from the fit are given in Table III.

discussed in Sec. III F. The observed value of μ_{eff} is slightly smaller than $2.54 \mu_B$ expected for a Ce³⁺ ion with $J = 5/2$. Further, as can be seen from Fig. 1(a), the inverse susceptibility also exhibits CW-type behavior between 30 and 135 K but with $\mu_{\text{eff}} = 2.56(2)\mu_B$ and $\theta_p = -31.8(3)$ K. The value of μ_{eff} estimated from the low-temperature CW behavior is similar to that expected for a Ce³⁺ ion. The large and negative value of θ_p indicates AFM interactions between the Ce ions. Such a large negative value of θ_p has also been observed in many Ce- and Yb-based HF systems and provides an estimate of the Kondo temperature through $T_K \sim |\theta_p|/4.5$ based on the s-d model.²⁶ Further within the symmetric Anderson model and local moment regime, Krishnamurthy *et al.* derived the relation $T_K = |\theta_p|/2$.²⁷ From our susceptibility data and using the s-d model, we obtain $T_K = 7$ K for CeRhGe₃. It should be noted that a similar method for estimation of T_K (estimated from $|\theta_p|/4$) has also been used for CeCuAl₃ and CeMg₃.^{28,29}

It is worth mentioning here that the single-crystal susceptibility of CeRhGe₃¹⁷ exhibits an anisotropy with the easy axis along $[100]$ [see Fig. 1(b)]. More specifically, the inverse susceptibility of a single crystal in both $[100]$ and $[001]$ directions exhibits CW behavior in two distinct temperature ranges, between 150 and 300 K and between 30 and 140 K [Fig. 1(b)]. The estimates of the effective paramagnetic moments and CW temperatures are $\mu_{\text{eff}} =$

$1.9 \mu_B$ ($1.9 \mu_B$) and $\theta_p = 48.3$ K (27.8 K) along $[100]$ ($[001]$) for the high-temperature range and $\mu_{\text{eff}} = 2.4 \mu_B$ ($2.3 \mu_B$) and $\theta_p = -19.8$ /K (-33.3 K) along $[100]$ ($[001]$) for the low-temperature range.¹⁷ We, therefore, see some commonality with our powder measurements, namely, the existence of two CW regions with ferromagnetic-type interactions and reduced moments in the high-temperature range, while the low-temperature range displays AFM-type interactions and effective moments closer to that expected for a Ce³⁺ ion. The presence of two different regions over which CW behavior is observed may suggest either the CEF parameters are temperature dependent, as observed in CeCuSi,³⁰ or the presence of anisotropic two-ion exchange interactions. Analysis of the single-crystal susceptibility [fit shown by solid lines in Fig. 1(b)] was carried out along with INS data and further discussion on this is given in Sec. III F. It is interesting to note that θ_p deduced from the single-crystal susceptibility is higher along c than along a .

C. Heat capacity

The heat capacity of CeRhGe₃ is shown in Fig. 2(a), together with that of the nonmagnetic phonon reference LaRhGe₃. The latter compound does not exhibit any anomaly or onset of SC down to 2 K. The low-temperature behavior of the heat capacity of LaRhGe₃ follows the form $C_p(T)/T = \gamma + \beta T^2$, indicating a phonon contribution to the heat capacity. Fitting the data below 6.0 K, we have estimated the value

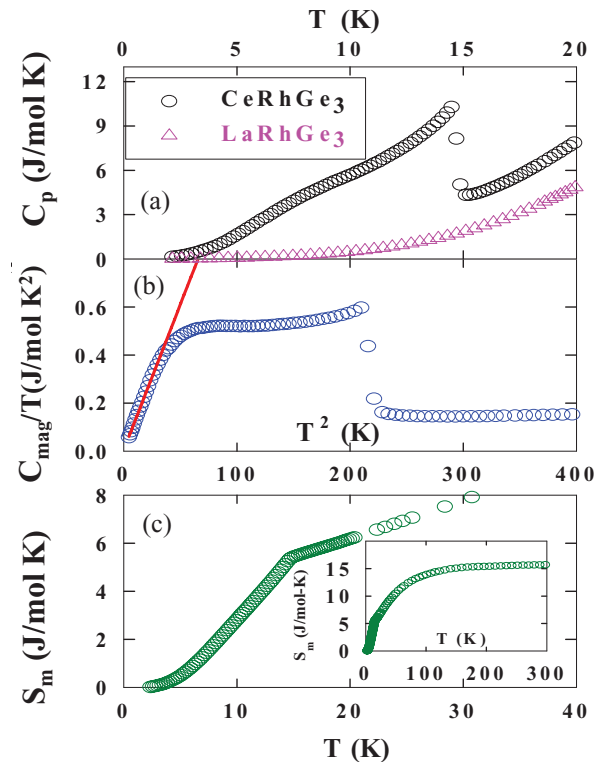


FIG. 2. (Color online) (a) Temperature dependence of the heat capacity of CeRhGe₃ and the phonon reference compound LaRhGe₃, (b) Estimated magnetic contribution of the heat capacity plotted as C_{mag}/T vs T^2 . The red solid line showing the fit (see text for details). (c) The temperature-dependent magnetic entropy estimated from the data in (b) and the inset shows the entropy up to 300 K.

of the Sommerfeld coefficient $\gamma \sim 5$ mJ/mole K² and $\beta = 0.415$ mJ/mole K⁴. This value of β gives a Debye temperature $\theta_D = 304$ K for LaRhGe₃. On the other hand, the heat capacity of CeRhGe₃ exhibits a sharp λ -type transition at 14 K [Fig. 2(a)] due to the onset of AFM ordering of the Ce moments in agreement with the magnetic susceptibility. Further, a very broad peak or anomaly in the heat capacity can be seen near 7 K in CeRhGe₃, which we suggest is due to either a change in the spin direction (or spin reorientation) of the Ce ions or a change in the propagation vector of the magnetic structure. The broad anomaly observed near 7 K is consistent with the peak observed in the magnetic susceptibility [see Fig. 1(a)], and the overall heat capacity behavior is in agreement with that reported by Muro *et al.*¹⁶ In order to investigate further the nature of these transitions, we also measured the heat capacity of CeRhGe₃ in an applied magnetic field of 9 T (data not shown here) and did not detect any change within the resolution of our measurements. This is in sharp contrast to the behavior observed in a polycrystalline sample of CeIrGe₃,³¹ where the application of a magnetic field moves the observed sharp low-temperature transition (at $T_{N2} = 4$ K) toward a higher temperature, while it does not affect the high-temperature transition at $T_{N1} = 8$ K. When the applied field reaches 7 T, the lower transition merges with the upper transition, and only one broad peak near 8 K can be seen in the heat capacity data.³¹ It is very interesting to note that the application of pressure and applied magnetic field has a similar effect on the two phase transitions in CeIrGe₃.^{17,31}

The magnetic (or 4f-electronic) contribution to the heat capacity of CeRhGe₃ was obtained by subtracting the lattice contribution from the measured heat capacity of LaRhGe₃ (i.e., by taking the difference $C_{\text{mag}}(T) = C_p(\text{CeRhGe}_3) - C_p(\text{LaRhGe}_3)$). The resulting curve is plotted in Fig. 2(b) as $C_{\text{mag}}(T)/T$ vs T^2 . A very sharp and well-pronounced transition at 14 K is clearly seen in the magnetic part of the heat capacity $C_{\text{mag}}(T)$, which further exhibits a drop near 7 K. Although the amplitude of the change $\Delta C_{\text{mag}}(T)$ in the magnetic heat capacity, near the 14 K jump is similar to that of the 7 K drop, the jump occurs over a much narrower temperature range than the drop. A broad peak in C_{mag} near 40 K (not shown) is attributed to a crystal electric field Schottky anomaly. The magnetic contribution to the entropy was obtained by integrating the C_{mag}/T vs T plot and is shown in Fig. 2(c). From the temperature dependence of the magnetic entropy, we find that S_{mag} attains a value of $\sim R \ln 2$ ($= 5.75$ J/mole K) at 16 K, suggesting a CEF-split doublet ground state in CeRhGe₃. Furthermore, the entropy recovery at 300 K is 15.67 J/mole K, which is close to $R \ln 6 = 14.87$ J/mole K, indicating that the overall CEF splitting is smaller than 300 K, in agreement with our INS study discussed in Sec. III F. We have made an estimate of the Kondo temperature T_K of CeRhGe₃ using the method described by Besnus *et al.*,³² where in the mean-field approach, the jump in the heat capacity ΔC_{mag} of a Kondo system is related to T_K/T_N . The estimate of T_K through this model agrees well with the experimental data on various Ce and Yb compounds. A correlation between the specific heat jump ΔC_{mag} at the magnetic ordering temperature and the ratio T_K/T_N for CeNi_xPt_{1-x} has been observed by Blanco *et al.*³³ Besnus *et al.* have given a universal plot of ΔC_{mag} vs T_K/T_N

for many Ce-based Kondo lattice systems. Using the observed value $\Delta C_{\text{mag}} = 6.44$ J/mole K of CeRhGe₃, we deduced $T_K/T_N = 0.8$ from this universal plot that gave $T_K = 11.8$ K for CeRhGe₃. This value of T_K is in reasonable agreement with that estimated from θ_p as well as that estimated from the quasielastic linewidth discussed in Sec. III F. The influence of the CEF above 20 K and the magnetic order below 14.5 K makes it difficult to estimate a reliable value of the Sommerfeld coefficient γ for CeRhGe₃. An attempt has been made to estimate γ from a linear fit to C/T vs T^2 plot above T_{N1} (between 15 to 20 K), yielding $\gamma \sim 99$ mJ/mole K². However, the C/T value at 2 K is only ~ 65 mJ/mole K²: this clearly reflects the influence of the CEF above the transition.

We now consider the spin-wave contribution to the heat capacity. From spin-wave theory, the temperature dependence of the magnetic specific heat for ferromagnets and antiferromagnets is proportional to $T^{3/2}$ and T^3 , respectively, for temperatures below the ordering temperature. The C_{mag}/T data of CeRhGe₃ were plotted as T^2 and also as $T^{0.5}$, and it was found that the T^3 (i.e., C_{mag}/T vs T^2 plot) dependence fits better [Fig. 2(b)] below T_{N2} , in agreement with the AFM ground state observed in CeRhGe₃. For a three-dimensional (3D) antiferromagnet with a linear spin-wave dispersion relation (below T_N), the coefficient of the T^3 term is given by $C_{\text{mag}} = (8\pi R/15)(T/\theta_c)^3$, where the parameter θ_c is of the order of T_N .³⁴ From the measured slope, we have estimated $\theta_c = 10.5$ K, which is indeed of the order of T_{N2} .

D. Muon spin relaxation

To shed light on the two phase transitions seen in the susceptibility and heat capacity, we have investigated the temperature dependence of the muon spin relaxation in zero field. For all temperatures above the magnetic ordering temperature, the muon spin relaxation spectra can be described by a simple exponential decay. However, on cooling below the ordering temperature ($T_{N1} = 14.5$ K), coherent frequency oscillations are observed in the μ SR spectra (Fig. 3). This immediately shows that the system has a long-range magnetically ordered ground state. The μ SR spectra are well described by three sinusoidal functions with a Gaussian envelope and an

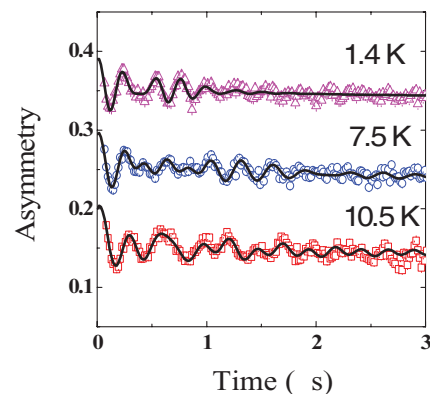


FIG. 3. (Color online) The time evolution of the muon spin relaxation for various temperatures (from top to bottom 1.4, 7.5, and 10.5 K). The line is least-squares fit to the data as described in the text. Note the plots at 7.5 and 1.4 K are shifted vertically (by 0.1) for clarity.

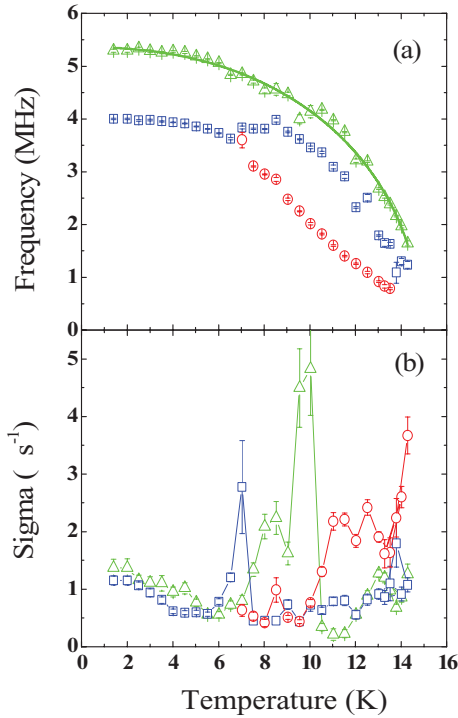


FIG. 4. (Color online) (a) The temperature dependence of the μ SR frequencies for CeRhGe₃. The line is a fit to the data as described in the text. (b) The temperature dependence of the Gaussian envelope σ for CeRhGe₃.

exponential decay (see Fig. 3) given by the functional form

$$G_z(t) = \left(\sum_{i=1}^3 A_i \cos(2\pi \nu_i t + \varphi) \exp\left(-\frac{(\sigma_i t)^2}{2}\right) \right) + A_0 \exp(-\lambda t) + A_{\text{background}}, \quad (2)$$

where A_i (A_0) is the initial asymmetry, σ_i is the Gaussian envelope to the oscillating function with a frequency of ν_i and a phase φ , λ is the muon spin relaxation rate, and $A_{\text{background}}$ is the background. Although one might expect the addition of three exponential relaxation functions, the data suggest that these values of λ would be so close that they can be fitted with one value.

Next, we consider the temperature dependence of the muon frequencies below T_{N1} shown in Fig. 4(a). There are three distinct frequencies for temperatures greater than ~ 7 K. However, as the temperature is reduced below 7 K, the middle frequency merges into the lowest frequency. Normally, three frequencies imply three distinct muon sites, but considering the crystal structure of CeRhGe₃, it is not obvious where these three sites could be. Interestingly, as discussed in Sec. III E, the magnetic structure of CeRhGe₃ is a spin-density wave type (see Fig. 6), and hence there are two types of crystallographic unit cells in the magnetic unit cell at 2 K. In one cell the Ce-layer (made up of four Ce atoms) has a z-component of the magnetic moment [see Fig. 6(a) bottom unit cell], while in the next cell this Ce layer does not have a z-component of the magnetic moment [zero moment, see Fig. 6(a) or smaller moment, see Fig. 6(b) second unit cell from the bottom]. This can explain the observed two frequencies below 7 K. Intriguingly, the presence

of the three frequencies above 7 K indicates that there is clearly a change in the magnetic structure with temperature, i.e., we have two different magnetic structures below and above 7 K. It suggests that we have a complex magnetic structure also above 7 K, which is likely to be incommensurate with the lattice.

The temperature dependence of the upper frequency, which is a measure of the internal field at the muon site, can be fitted with

$$\nu(T) = \nu(0) \left(1 - \left(\frac{T}{T_N} \right)^\alpha \right)^\beta. \quad (3)$$

The results of the fit gives $\nu(0) = 5.40(6)$ MHz, $T_N = 14.8(2)$ K, $\alpha = 2.3(2)$ and $\beta = 0.48(7)$, which being close to $\frac{1}{2}$ indicates that CeRhGe₃ can be thought of as a mean-field magnet. The value of α , which is much greater than 1, implies that there are complex interactions between the moments as inferred through our neutron diffraction study in Sec. III E that reveals a complex spin arrangement in the magnetic unit cell.

Finally, the temperature dependence of σ is shown in Fig. 4(b). Near the temperature at which we observe the three frequencies merging into two, we also see a peak in $\sigma(T)$. Further, the $\sigma(T)$ also exhibits a peak near 10 K, where the highest frequency exhibits a weak anomaly. The peaks in the $\sigma(T)$ indicate an increase in the distribution of the internal fields at the muon site(s) around the main field. Each peak shows that either an increase in the spin fluctuations or a very small change in the magnetic structure occurs, possibly a small

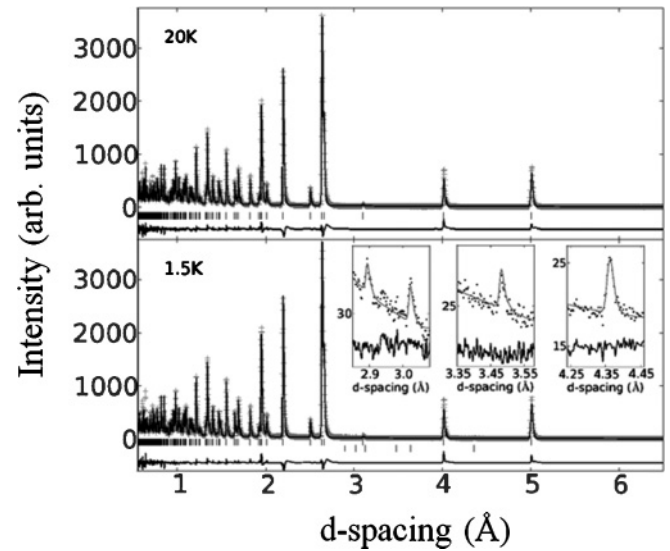


FIG. 5. Rietveld refinement of the neutron powder diffraction pattern (obtained from WISH) of CeRhGe₃ at 20 K (top) and 1.5 K (bottom). The data are shown as grey crosses, and the result of the refinement as a solid (black) line. The upper and lower rows of tick marks indicate the position of nuclear and magnetic Bragg peaks, respectively. The difference curve between the experimental data and the refinement is shown at the bottom of both figures. The insets in the bottom figure display an enlarged region of the diffraction pattern near the magnetic Bragg peaks along with the refinement.

change in the modulation wave vector (\mathbf{k}) of the magnetic moment, within the μ SR time window.

E. Magnetic neutron diffraction

Figure 5 shows the neutron diffraction data collected at 20 and 1.5 K for the 90° detector bank on WISH. The refinement (weighted profile factor $R_{wp} = 5.34\%$) at 20 K confirms that CeRhGe_3 crystallizes in the tetragonal space group $I4mm$, in agreement with both our GEM data and the previous report,¹⁷ with $a = 4.385(2)$ and $c = 10.015(3)$ Å at 20 K. The presence of a small impurity, which could not be indexed by any of the binary and tertiary compounds of Ce-Rh-Ge reported in the ICSD database³⁵ was also detected. At 1.5 K, the diffraction data reveals the presence of four clear additional peaks at 4.36, 3.48, 3.02, and 2.89 Å (insets of Fig. 5), and two possible very weak peaks at 2.17 and 1.8 Å that are better resolved in the higher-angle detector banks (not shown). The intensity of these peaks decreases with the modulus of the scattering vector (Q) as expected from the Q -dependence of the magnetic form factor. It is to be noted that only Ce atoms have magnetic moment (Rh and Ge atoms do not carry magnetic moment), and there is only one crystallographic site for the Ce atoms per magnetic unit cell and other Ce atoms are related by the symmetry operators. Further evidence of the magnetic origin of these extra peaks is provided by the reduction of the background at low Q . This is the expected behavior at a magnetic transition where the intensity of the paramagnetic scattering is transferred to the Bragg scattering. Initially, an automatic indexing procedure using a grid search was used to determine the periodicity of the magnetic structure. To confine the wave vector search, all the special points and lines of the Brillouin zone generated by the program ISOTROPY³⁶ were tested. The best solution, accounting for all the observed peaks, was found along the line $LD(0,0,2a)$ for the wave vector $\mathbf{k} = (0,0,3/4)$. This wave vector corresponds to $(3/8, 3/8, -3/8)$ in the primitive cell and can be labeled k_{10} in Kovalev's³⁷ notation. At this point, it is worth mentioning that the same wave vector $\mathbf{k} = (0,0,3/4)$ has also been observed in a recent single-crystal neutron diffraction study by Kaneko *et al.*²¹ on isostructural CeCoGe_3 . However, these authors also reveal the existence of another dominant wave vector $\mathbf{k}' = (0,0,1/2)$, not detected in our CeRhGe_3 sample, with intensities associated with \mathbf{k}' being a factor 7 higher than those associated with $\mathbf{k} = (0,0,3/4)$. None of the other special points of the Brillouin zone, nor a full 3D grid search, gave a better R-factor. Symmetry analysis indicates that there are four, one-dimensional (1D) representations, labeled Γ_1 to Γ_4 and one, two-dimensional (2D) representation Γ_5 in the little group (group of the propagation vector), although only Γ_2 and Γ_5 enter the decomposition of $\Gamma_{\text{mag}} = \Gamma_2 + \Gamma_5$. Γ_2 and Γ_5 correspond to an ordering of the Ce sites along the c axis and in the ab -plane, respectively. A good fit to the data (magnetic Bragg factor $R_B = 11.9\%$) was obtained using FullProf³⁸ with the representation Γ_2 . Attempts at refinements using Γ_5 gave an R_B a factor 6 to 7 times higher with the peaks at 2.89, 3.02, and 4.36 Å being fairly well accounted for, but the one at 3.48 Å overestimated. Using Γ_5 also introduces some intensity in the $(002) + \mathbf{k}$ peak at 3.64 Å, which is not very clear in the 90° bank data and

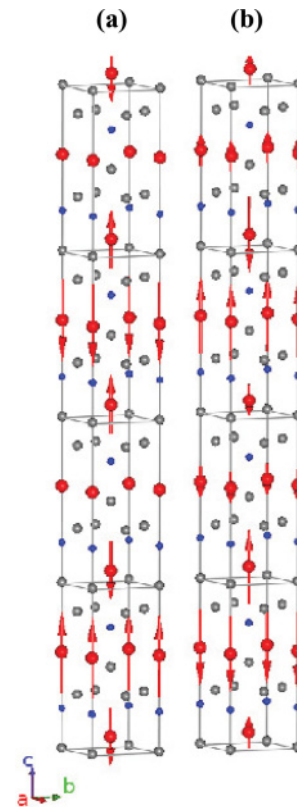


FIG. 6. (Color online) Magnetic unit cell of CeRhGe_3 (a) with zero value of the global phase and (b) with the global phase of $9/16 * \pi \text{ mod } (\pi/4)$. The large red (dark gray), small blue (medium gray), and large gray symbols indicate Ce, Rh, and Ge atoms, respectively. The red arrows show the direction and size of the Ce magnetic moment.

also in the $(002)\text{-k}$ peak at 8.02 Å is not visible in the lower-angle diffraction bank. The refined moment, using the Γ_2 representation, was $0.45(9) \mu_B$, and the magnetic structure is depicted in Fig. 6. It should be noted that the moment direction obtained from our neutron diffraction study at 1.5 K is along the c axis, which is in agreement with the drop in the c axis susceptibility.¹⁷ On the other hand, the easy axis (i.e., the axis with the highest magnetization in applied magnetic field) observed in the antiferromagnetic (AFM) state from the magnetization data at 2 K is the a axis (see Fig. 5 in Ref. 17) and not the c axis. Further, the observed moment along the c axis in CeRhGe_3 is also in contrast with the highest susceptibility direction, which is the a axis for $T > T_N$.¹⁷

It should be noted that neutron diffraction patterns obtained from a given magnetic structure are identical when a global phase Φ is added. Figure 6(a) shows the magnetic structure obtained for a Φ equals to zero and includes some Ce sites having zero moment. In the present case, no suitable global phase that would result in a constant moment solution could be found but a special case when the global phase = $9/16 * \pi \text{ mod } (\pi/4) = 0.5625 * \pi \text{ mod } (\pi/4)$ [see Fig. 6(b)] for which the values of the moments on the first Ce1 site at $(0\ 0\ 0.579)$ are equal the ones on the second Ce2 site at $(0.5\ 0.5\ 0.579)$ (still the moments are not constant on either of the sites and are along the c axis), i.e., Ce1 atoms translated by $(0\ 0\ n)$

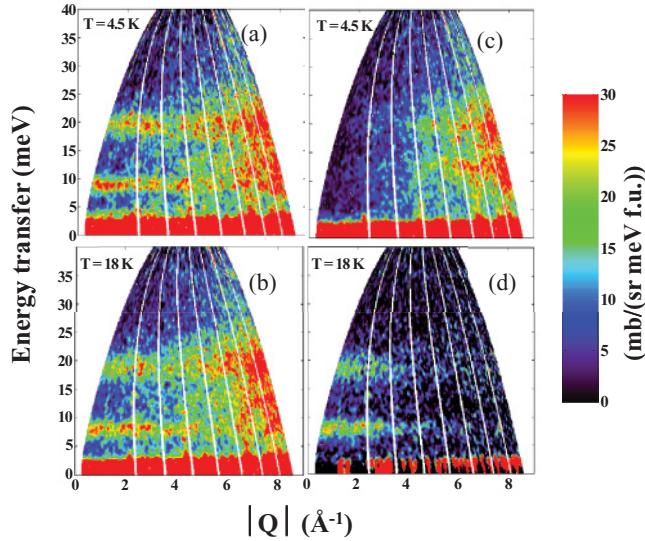


FIG. 7. (Color online) Contour plots of the inelastic scattering (in mb/sr meV f.u.) with energy transfer (E) vs wave vector transfer (Q), (a) and (b) of CeRhGe₃ at 4.5 and 18 K and (c) of LaRhGe₃ at 4.5 K. (d) The magnetic scattering of CeRhGe₃ at 18 K estimated by subtracting the data of LaRhGe₃.

($n = 0, 1, 2, 3$) have a moment of $\mu_z = -0.42, -0.17, 0.42, 0.17 \mu_B$, respectively, and Ce2 atoms translated by $(00n)$ have a moment of $\mu_z = 0.17, 0.42, -0.17, -0.42 \mu_B$, respectively. As we are unable to determine the muon site in CeRhGe₃ from our present μ SR study on the polycrystalline sample, single-crystal nuclear magnetic resonance (NMR) and μ SR studies should prove very useful to determine the unique value of the global phase of the magnetic structure of CeRhGe₃.

F. Inelastic neutron scattering

1. High-energy INS study to investigate CEF excitations

Figures 7(a) and 7(b) show contour plots of the INS, measured on the MARI spectrometer with $E_i = 45$ meV, from CeRhGe₃ at 4.5 and 18 K and also from the phonon reference compound LaRhGe₃ at 4.5 K [Fig. 7(c)]. In the paramagnetic state (at 18 K $> T_N$) at low scattering vector (Q), two inelastic excitations near 7.5 and 18 meV can be clearly seen in CeRhGe₃. Similar scattering has also been observed at 4.5 and 10 K, which indicates that scattering from the spin waves in the ordered state of CeRhGe₃ is at very low energy below 3.5 meV and is confirmed by our low-energy study discussed in Sec. III F2. In contrast the scattering in LaRhGe₃ is very weak at low Q [Fig. 7(c)] but exhibits at high Q the same two peak structure, as seen in CeRhGe₃, with nearly similar intensity. This shows the presence of two magnetic excitations in CeRhGe₃ having similar energies to the phonon peaks. It is interesting to mention that despite having similar energy scales, the magnetic and phonon excitations appear uncoupled and do not show any CEF-phonon coupling effect, which would give rise to an extra magnetic-CEF peak over the number expected from Kramers theorem.³⁹ According to this theorem, one cannot observe more than two CEF excitations from the $J = 5/2$ ground state of Ce³⁺ ion in any Ce compound in the paramagnetic state. Such CEF-phonon coupling has been

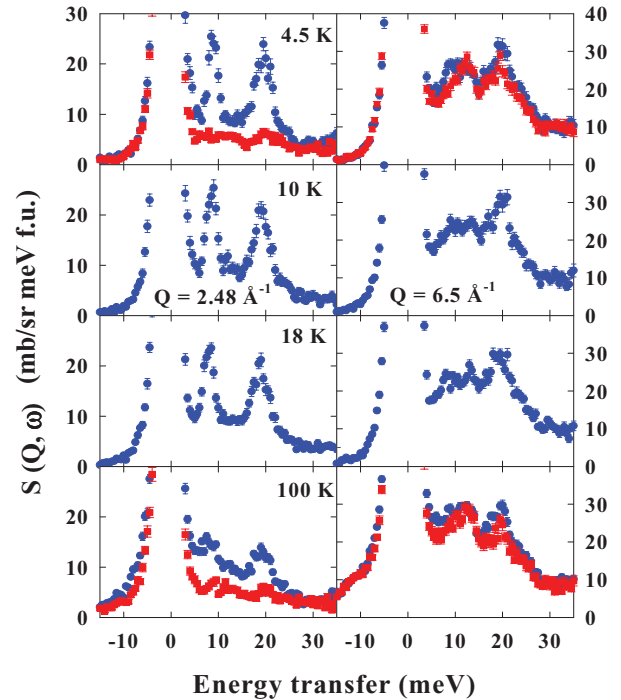


FIG. 8. (Color online) The Q -integrated 1D cuts of the total scattering from CeRhGe₃ (blue circles) and LaRhGe₃ (red squares) at low $Q = 0$ to 4 \AA^{-1} (left side) and at high $Q = 5$ to 9 \AA^{-1} (right side).

detected in CeCuAl₃, which has a similar crystal structure to CeRhGe₃, where three CEF excitations from the ground state are clearly observed.⁴⁰

The scattering at the highest Q is comparable for CeRhGe₃ and LaRhGe₃, which indicates similar phonon contributions in these compounds. This can be seen clearly in the 1D cuts made from the 2D color plots at low Q from 0 to 4 \AA^{-1} and at high Q from 5 to 9 \AA^{-1} (see Fig. 8) at 4.5, 10, 18, and 100 K. At 4.5 and 100 K the phonon contributions at high Q are the same in both compounds, but at low Q , the magnetic scattering is strong and with a small phonon contribution in CeRhGe₃. The magnetic scattering has been deduced by subtracting the observed scattering for LaRhGe₃ from that of CeRhGe₃, i.e., $S_M(Q, \omega) = S(Q, \omega, \text{CeRhGe}_3) - S(Q, \omega, \text{LaRhGe}_3)$. This is shown in Fig. 7(d), which clearly reveals the presence of two excitations. The Q -integrated (from 0 to 4 \AA^{-1}) 1D cuts of the magnetic scattering are shown in Fig. 9 at all four temperatures. Figure 9(c) shows two well-resolved inelastic excitations at 18 K (i.e., above T_N), which we attribute to the CEF excitations from the ground-state CEF level to two excited CEF levels: the $J = 5/2$ ground-state multiplet of the Ce³⁺ ion in CeRhGe₃ splits into three CEF doublets and hence gives two CEF excitations. To check the Q -dependence of the intensity of these excitations, we have plotted the energy integrated intensity between 6 and 9 meV and between 16 and 21 meV, as a function of Q in Fig. 10. The intensity of both excitations decreases with increasing Q , following the square of the magnetic form factor $F^2(Q)$ for a Ce³⁺ ion. A similar Q -dependence was also observed at 4.5 K. As

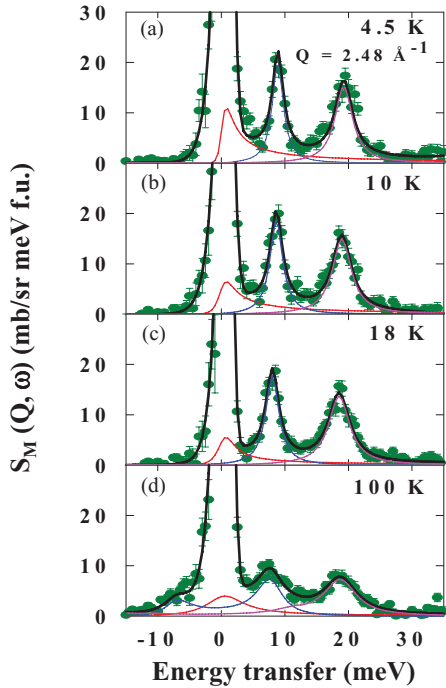


FIG. 9. (Color online) The estimated magnetic scattering from CeRhGe₃ at low $Q = 0$ to 4 \AA^{-1} at 4.5, 10, 18, and 100 K. The thick solid lines represent the fit (based on the CEF model), and thin solid lines show the components of the fit (see text).

expected, the phonon scattering increases with Q and follows a Q^2 dependence.

On further decreasing the temperature to 10 and 4.5 K, a small increase in the energy of the CEF excitations can be noticed compared to the values at 18 K, which is due to either the presence of a molecular field in the magnetically ordered state or the possibility of magnetostriction below T_N . At 18 K the linewidth of the 7.5 meV excitation is $1.4(2)$ meV, while that of 18 meV excitation is $2.2(3)$ meV for CeRhGe₃. This latter value is significantly greater than the instrumental resolution of 1.0 meV for this energy transfer. Considering that CeRhGe₃ is a crystallographically ordered system with one Ce site, the larger linewidth of the higher-energy excitation, compared to the resolution, may be attributed to the presence of a strong hybridization between the wave functions of this particular CEF level with the conduction electrons, indicating the presence of anisotropic hybridization. It is interesting to compare the energy scale and linewidths of the CEF excitations in CeRhGe₃, with those of CeRhSi₃, which has same crystal structure but shows pressure-induced SC with $T_{sc} = 1$ K at 2.5 GPa. The INS study on CeRhSi₃ shows two CEF excitations at 19 and 36 meV with linewidths of $3.9(2)$ and $9.2(4)$ meV, respectively.⁷ The overall CEF splitting and linewidths in CeRhSi₃ are higher than those observed in CeRhGe₃, which indicates stronger hybridization for the Si compound. The weaker hybridization observed in CeRhGe₃ compared with that in CeRhSi₃ through our INS study is also in agreement with that proposed by Kawai *et al.*¹⁷ based on the molar volume vs T_N and molar volume vs γ plots. In these plots, CeRhSi₃ is close to a QCP, while CeRhGe₃ is well below the QCP (on the localized limit side with

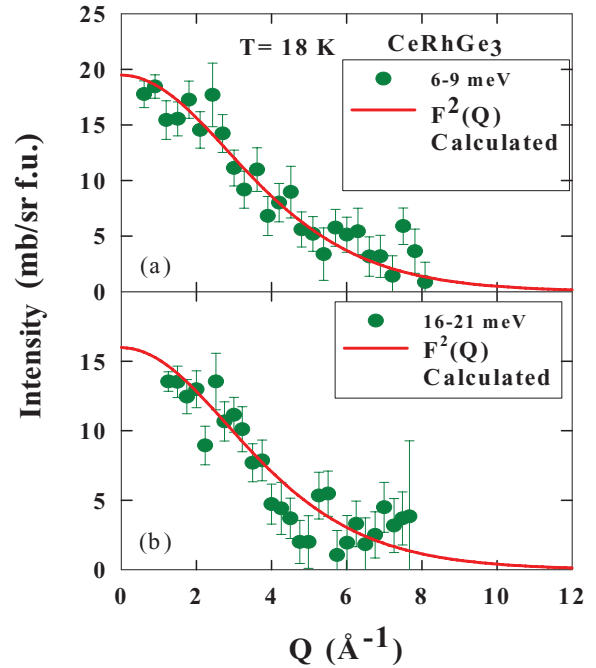


FIG. 10. (Color online) The energy integrated [from 6–9 meV (a) and 16–21 meV (b)] Q -dependent magnetic scattering from CeRhGe₃ at 18 K. The solid line represents the Ce³⁺ magnetic form factor squared calculated from P. J. Brown's tables (Ref. 53).

smaller hybridization). The observation of well-resolved CEF excitations in CeRhGe₃ supports Kawai's analysis.

We now proceed with a more detailed analysis of the observed CEF excitations. In the tetragonal point symmetry, 4 mm (C_{4v}), at the Ce site, the CEF Hamiltonian can be represented as follows:

$$H_{CF} = B_2^0 O_2^0 + B_4^0 O_4^0 + B_4^4 O_4^4, \quad (4)$$

where B_n^m are the CEF parameters to be determined from the experimental data and O_n^m are Stevens operator equivalents obtained using the angular momentum operators.⁴¹ The value of B_2^0 can be accurately determined using the high-temperature expansion of the magnetic susceptibility,⁴² which gives B_2^0 in terms of the CW temperatures, θ_{ab} for an applied field in the ab -plane and θ_c for an applied field parallel to c axis:

$$B_2^0 = \frac{10}{3}(\theta_{ab} - \theta_c)/(2J - 1)(2J + 3). \quad (5)$$

Note that Eq. (5) is valid for isotropic exchange interactions (i.e., isotropic molecular field parameters). Using the values of θ_{ab} and θ_c from the single-crystal susceptibility,¹⁷ we find $B_2^0 = 0.206$ meV from the high-temperature CW behavior between 150 and 300 K and $B_2^0 = 0.135$ meV from the low-temperature CW behavior between 30 and 135 K. In our subsequent analysis, we first kept the value of B_2^0 (both values were tried) from this estimate fixed and then varied B_4^0 and B_4^4 . Although we could get a good fit to the neutron data at 18 K and 100 K with these values of B_2^0 , the estimated CEF parameters did not give a good fit to the single-crystal susceptibility data for the entire temperature range from $T > T_N$ to 300 K. Hence in the final fit, we also allowed B_2^0 to vary and fitted the neutron scattering data (at each temperature) together with the single-crystal susceptibility. The values of the parameters

TABLE III. CEF parameters B_n^m , molecular field parameters (λ^ξ , $\xi = \perp$ and \parallel to c axis) and temperature-independent constant susceptibility (χ_0^ξ), calculated CEF ground-state magnetic moments ($\langle\mu_x\rangle$ and $\langle\mu_z\rangle$ calculated from the CEF ground-state wave functions) and the calculated energy (ΔE_1 and ΔE_2 are the energy of first- and second-excited CEF levels) of the CEF levels. The parameters were estimated by simultaneous fit to the inelastic neutron data and the single-crystal susceptibility at each temperature.

	4.5 K	10 K	18 K	100 K
B_2^0 (meV)	0.381	0.385	0.393	0.408
B_4^0 (meV)	0.021	0.021	0.018	0.018
B_4^4 (meV)	0.308	0.302	0.294	0.301
λ^\perp (mole/emu)	-17.286	-17.586	-19.787	-19.919
λ^\parallel (mole/emu)	10.939	8.671	8.079	10.396
χ_0^\perp ($\times 10^{-3}$ emu/mol)	-1.086	-1.090	-1.091	-1.094
χ_0^\parallel ($\times 10^{-3}$ emu/mol)	-0.964	-0.948	-0.940	-0.931
$\langle\mu_x\rangle$ (μ_B)	0.83	0.83	0.83	0.83
$\langle\mu_z\rangle$ (μ_B)	0.43	0.44	0.43	0.42
ΔE_1 (meV)	8.73	8.53	7.66	7.62
ΔE_2 (meV)	19.10	18.84	18.20	18.58

obtained from this procedure are given in Table III, and the quality of the fit to the measured data can be seen in Fig. 1(b) for the susceptibility and Fig. 9 for the neutron scattering data. The estimated value of B_2^0 is higher than that found from the susceptibility, while B_4^0 is very weak, and B_4^4 is the second largest parameter. The CEF wave functions obtained from the simultaneous fit to 18 K INS and the susceptibility data are given by

$$\begin{aligned}\Psi_1^\pm &= (0.8658)|\pm \frac{3}{2}\rangle - (0.5004)|\mp \frac{5}{2}\rangle \\ \Psi_2^\pm &= |\pm \frac{1}{2}\rangle \\ \Psi_3^\pm &= (0.8658)|\pm \frac{5}{2}\rangle + (0.5004)|\mp \frac{3}{2}\rangle\end{aligned}\quad (6)$$

They correspond to the energy eigenvalues of 0, 7.7, and 18.2 meV. We can now estimate the value of the ground state magnetic moment using the above wave functions and the following formulae:

$$\begin{aligned}\langle\mu_x\rangle &= \langle\Psi_1^\pm|\frac{gJ}{2}(J^+ + J^-)|\Psi_1^\mp\rangle \\ \langle\mu_z\rangle &= \langle\Psi_1^\pm|gJ(J_Z)|\Psi_1^\pm\rangle.\end{aligned}\quad (7)$$

The estimated magnetic moments $\langle\mu_x\rangle$ and $\langle\mu_z\rangle$ are given in Table III. The value of $\langle\mu_z\rangle$ estimated here is in excellent agreement with that obtained from the neutron diffraction study. However, on the basis of the CEF model one would expect the ordered state Ce moment in the ab-plane and not along the c axis. Now we consider the direction of the moment. According to the single-crystal susceptibility data in the paramagnetic state and also from the analysis of our INS data, B_2^0 is positive. If we consider the magnetocrystalline anisotropy energy in terms of the second-order CEF parameter, which is given, in the lowest order term in the energy by⁴³

$$E_a = K_1 \sin^2(\theta). \quad (8)$$

Here θ is the angle between the moment direction and the $[0\ 0\ 1]$ axis. If $K_1 > 0$, the directions of lowest energy are the $\pm z$ directions, and z axis is the easy axis. If $K_1 < 0$,

there is an easy plane perpendicular to the symmetry axis (the basal plane of the crystal). Hence from the sign of B_2^0 one can predict the magnetization direction. For Ce^{3+} systems a positive B_2^0 ($K_1 < 0$) minimizes E_a when the angle θ is 90° , i.e., moment in the ab-plane, while for negative B_2^0 ($K_1 > 0$), E_a will be minimum for the moment along the c axis. In $CeRhGe_3$ we have a positive B_2^0 , which predicts that the moment should be in the ab-plane. But the observed moment at 1.5 K is along the c axis. This remains the case even when we include the B_4^4 term (the weak B_4^4 can be neglected) in the anisotropy energy. This suggests that an anisotropy other than the magnetocrystalline anisotropy is playing an important role in determining the direction of the moments in $CeRhGe_3$. We attribute this anomaly of the magnetic moment direction in $CeRhGe_3$ to the presence of two-ion anisotropic magnetic exchange interactions between the Ce ions, along the c axis and in the ab-plane.⁴⁴ This interpretation has been supported through the estimated anisotropic molecular field parameters (see Table III) from the susceptibility analysis, negative sign for λ^\perp and positive sign for λ^\parallel . This type of behavior has also been observed in the ferromagnetic compound $CeAgSb_2$, where the CEF predicts moments in the ab-plane, but the experimental moment direction is along the c axis.^{45,46} As this is a ferromagnet, the change in the easy axis direction can be clearly seen in the temperature dependence of the single-crystal susceptibility, which shows a crossover below the magnetic ordering temperature.⁴⁶ This is not the case for antiferromagnetically ordered systems, where no such crossover is seen. Instead, the easy magnetization axis direction can be determined by examining the temperature-dependent susceptibility, which decreases below T_N for the direction parallel to the sublattice magnetization,⁴⁷ while the perpendicular susceptibility remains almost constant (or is weakly temperature dependent) below T_N . This is the case for $CeRhGe_3$ and very similar behavior has been observed in $CeRhIn_5$ and $CePt_3Si$.⁴⁸⁻⁵¹ The neutron diffraction study of $CeRhIn_5$ reveals that the orientation of the magnetic moments is in the ab-plane, while the susceptibility shows an easy

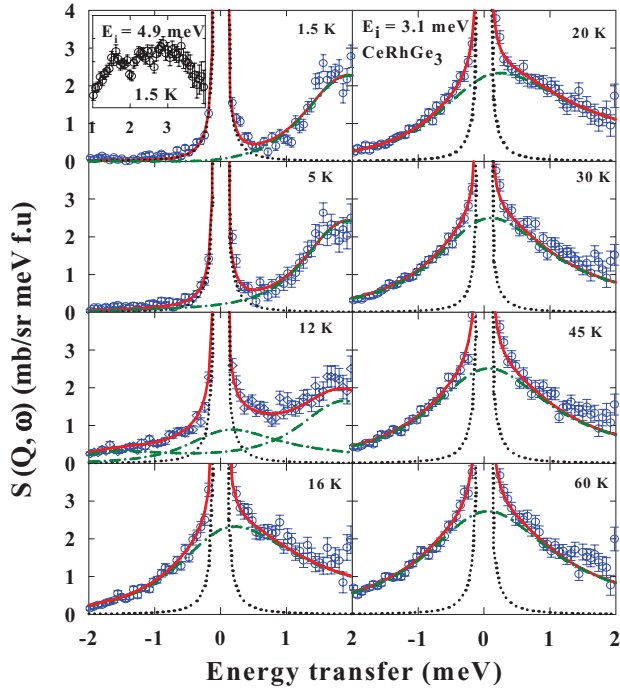


FIG. 11. (Color online) The observed scattering between 10° and 115° scattering angles for CeRhGe_3 at various temperatures measured with $E_i = 3.1$ meV. The data were fitted using a Lorentzian line-shape function for the quasielastic and inelastic components. The solid line shows the fit and dashed (elastic peak) and dashed-dotted (quasielastic and inelastic peaks) lines represent the components of the fit. The inset (top left) shows the data measured with $E_i = 4.9$ meV at 1.5 K.

magnetization direction along the c axis.^{48,49} For CePt_3Si , the single-crystal susceptibility and CEF parameters obtained from the INS analysis predict an easy magnetization along the c axis,⁵⁰ but the neutron diffraction data show that the moments are in the ab -plane.⁵¹ Recently, similar behavior has been observed in $\text{CeRu}_2\text{Al}_{10}$.⁵² These observations suggest that a systematic study, both theoretical and experimental, on Ce-based systems, is highly necessary to obtain a complete understanding of the role of anisotropic exchange and single-ion crystal-field anisotropies on the magnetism of these strongly correlated electron systems.

2. Low-energy INS study

To elucidate the nature of the low-energy excitations, Kondo temperature and spin-wave energy scale, we have also performed low-energy INS measurements on CeRhGe_3 , and the results are shown in Fig. 11. The data are plotted as the total scattering from all the detectors for scattering angles from 10° to 115° . At 1.5 K the scattering was mainly dominant below 0.7 \AA^{-1} and above 1 meV, which confirmed its spin-wave origin. In order to cover the high-energy part of the spin waves, we also carried out measurements with higher $E_i = 4.9$ meV at 1.5 K, and the integrated scattering is plotted in the inset of Fig. 11 (top left side). It is clear from the inset that the scattering intensity has a peak near 3 meV with a possibility of another peak near 2 meV. This suggests the presence of two (or more) possible spin-wave modes. At 5 K the spin-wave scattering is almost identical to

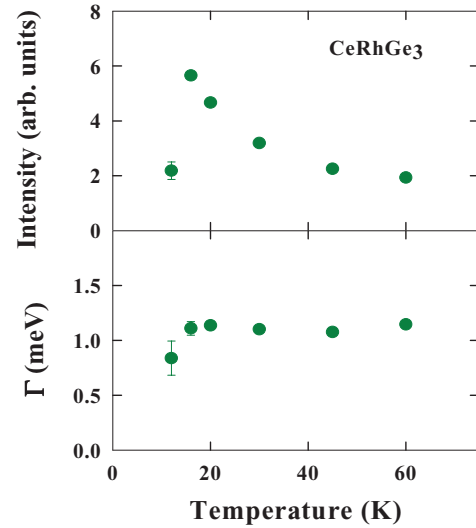


FIG. 12. (Color online) Temperature dependence of the intensity (top) and linewidth (bottom) of the quasielastic line estimated from the fit (see text).

that observed at 1.5 K, but at 12 K, we can see an increase in the quasielastic scattering and a slight decrease in the spin-wave intensity. Considering the numbers of spin-wave modes equal to the numbers of magnetic atoms per magnetic unit cell, one expects two degenerate spin-wave modes for a body-centered tetragonal collinear AFM (type-I AFM) structure (with equal moments on both Ce atoms, two Ce atoms per magnetic unit cell) for an isotropic magnetic exchange. It is to be noted that an anisotropic magnetic exchange could lift the degeneracy of the spin-wave modes. Furthermore the spin density wave-type AFM structure of CeRhGe_3 , having six Ce atoms carrying magnetic moment per magnetic cell (four crystallographic cells), will give three degenerate spin-wave modes for an isotropic magnetic exchange and six modes for an anisotropic magnetic exchange. The observed two (or possibly more) peaks in CeRhGe_3 can be attributed to the powder averaging from the ab -plane and along the c axis. However, to find out the real origin of the observed two (or more) peaks in the spin-wave scattering a detailed spin-wave investigation on a single crystal of CeRhGe_3 is highly desirable.

In the paramagnetic state, between 16 and 60 K, a clear sign of quasielastic scattering has been observed (Fig. 11). From the data it is evident that the quasielastic linewidth (Γ , HWHM) does not show any dramatic change with temperature. Further, the Q -dependent intensity in the paramagnetic state follows the square of the Ce^{3+} form factor. To estimate the temperature dependence of Γ , we have analyzed the data using a Lorentzian line-shape function convoluted with the elastic line resolution function. The elastic line resolution was modeled with Gaussian and Lorentzian functions and their parameters were estimated (and kept fixed in the analysis of Γ) from the vanadium run under the same conditions. The estimated intensity and linewidth of the quasielastic scattering are plotted in Fig. 12. This shows that $\Gamma(T)$ is nearly temperature independent between 16 and 60 K and decreases slightly at 12 K. On the other hand, the intensity increases with decreasing temperature and drops at 12 K (Fig. 12). As at 5 and

1.5 K, the quasielastic scattering was absent, and the data were fitted with an inelastic peak with fixed position at 1.8 meV. By extrapolating $\Gamma(T)$ to $T = 0$ K, which gives the measure of T_K , we have estimated $T_K = 12.6(3)$ K for CeRhGe_3 , which is in excellent agreement with that estimated using the heat capacity data in Sec. III C.

IV. CONCLUSIONS

We have used magnetic susceptibility, heat capacity, neutron powder diffraction, INS, and muon spin relaxation studies to probe the magnetic ground state of CeRhGe_3 . We have observed two magnetic transitions in the heat capacity and susceptibility near 7 and 14.5 K. Our neutron diffraction measurements at 1.5 K reveal a spin-density-wave-type magnetic ordering with propagation vector $\mathbf{k} = (0\ 0\ \frac{3}{4})$ and a moment of $0.45(9)\ \mu_B$ along the c axis. The muon spin relaxation (μSR) measurements reveal the presence of internal fields below 14.5 K, and we have observed three clear frequencies in the μSR spectra in the temperature range of 7 to 14.5 K, which then merge into two frequencies as the temperature is decreased below 7 K. We attribute these three frequencies to a complex magnetic structure. At 1.5 K, we observed only two frequencies, in agreement with the neutron diffraction data. Both studies confirm that the ground state of CeRhGe_3 exhibits long-range magnetic order. From the temperature dependence of the μSR frequencies, we have estimated the value of the critical exponent $\beta \sim 0.5$. Further, our high-energy INS study at 18 K reveals the presence of two well-defined

CEF excitations. The observed direction of the moment along the c axis does not agree with that predicted by the single-ion CEF anisotropy estimated from the analysis of the INS data. This indicates that the direction of the moment is governed by some other anisotropies, probably two-ion anisotropic magnetic exchange as evident from the anisotropic molecular field parameters. A similar behavior has been observed in many other Ce-based HF systems and calls for more detailed experimental and theoretical investigations in these compounds. Our low-energy INS study reveals spin waves below T_N and well-defined quasielastic scattering above T_N . Finally, the comparison of the linewidth of the CEF excitations of CeRhGe_3 and CeRhSi_3 indicates a stronger hybridization in the latter case, which becomes a superconductor at 2 GPa, while no SC has been observed in CeRhGe_3 up to 8 GPa. This suggests that the role of the hybridization could be critical for the SC in CeTX_3 compounds. This is further supported through an onset of SC at ambient pressure in CeCoSi_3 , which has strongest hybridization among CeTX_3 systems.

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